## WHAT IS CLAIMED IS:

1. A method of accelerating wound healing, comprising applying to a wound an effective amount to accelerate wound healing of at least one peptide other than Angiotensin II, said peptide consisting essentially of at least three contiguous amino acids and having a sequence corresponding to a subsequence of groups R<sup>1</sup>-R<sup>8</sup> in general formula I:

 $R^1 - R^2 - R^3 - R^4 - R^5 - R^6 - R^7 - R^8$  (SEQ ID NO:15), wherein

R<sup>1</sup> is selected from the group consisting of Asp, Glu, Asn, Acpc, Ala, Me<sup>2</sup>Gly, Pro, Bet, Glu(NH<sub>2</sub>), Gly, Asp(NH<sub>2</sub>) and Suc;

R<sup>2</sup> is selected from the group consisting of Arg, Lys, Ala, Orn, Ser(Ac), Sar, D-Arg and D-Lys;

R<sup>3</sup> is selected from the group consisting of Val, Ala, Leu, Ile, Gly, Pro, Aib, Acpc and Tyr;

R<sup>4</sup> is selected from the group consisting of Tyr, Thr, Ser and azaTyr;

R<sup>5</sup> is selected from the group consisting of Ile, Ala, Leu, Val and Gly;

R<sup>6</sup> is His or Arg;

R<sup>7</sup> is Pro or Ala; and

R<sup>8</sup> is selected from the group consisting of Phe, Phe(Br), Ile and Tyr, excluding sequences wherein R<sup>4</sup> is an amino terminal Tyr group of the peptide.

- 2. A method according to Claim 1, wherein the peptide is administered in matrical or micellar solution.
- 3. A method according to Claim 1, wherein the peptide is administered in an amount of at least 0.1 ng per kg body weight in a suitable carrier or diluent.
- 4. A method according to Claim 3, wherein the carrier or diluent is selected from the group consisting of carboxymethyl cellulose preparations, crystalloid preparations, viscoelastics, polyethylene glycols and polypropylene glycols.
- 5. A method according to Claim 1, wherein the peptide is administered in conjunction with a wound dressing.
- 6. A method according to Claim 1, wherein the peptide has a sequence consisting essentially of  $R^1 R^2 R^3$  in general formula I.

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- 7. A method according to Claim 6, wherein the peptide has a sequence consisting essentially of Asp Arg Val (SEQ ID NO:11).
- 8. A method according to Claim 1, wherein the peptide has a sequence consisting essentially of  $R^1 R^2 R^3 R^4$  in general formula I.
- 9. A method according to Claim 8, wherein the peptide has a sequence consisting essentially of Asp Arg Val Tyr (SEQ ID NO:10).

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- 10. A method according to Claim 1, wherein the peptide has a sequence consisting essentially of  $R^1 R^2 R^3 R^4 R^5$  in general formula I.
- 11. A method according to Claim 10, wherein the peptide has a sequence consisting essentially of Asp Arg Val Tyr Ile (SEQ ID NO:9).
- 12. A method according to Claim 1, wherein the peptide has a sequence consisting essentially of  $R^1 R^2 R^3 R^4 R^5 R^6$  in general formula I.
- 13. A method according to Claim 12, wherein the peptide has a sequence consisting essentially of Asp Arg Val Tyr Ile His (SEQ ID NO:8).
- 14. A method according to Claim 1, wherein the peptide has a sequence consisting essentially of  $R^3 R^4 R^5 R^6 R^7$  in general formula I.
- 15. A method according to Claim 14, wherein the peptide has a sequence consisting essentially of Val Tyr Ile His Pro (SEQ ID NO:6).
- 16. A method according to Claim 1, wherein the peptide has a sequence consisting essentially of  $R^2 R^3 R^4 R^5 R^6 R^7$  in general formula I.
- 17. A method according to Claim 16, wherein the peptide has a sequence consisting essentially of Arg Val Tyr Ile His Pro (SEQ ID NO:5).
- 18. A method according to Claim 1, wherein the peptide has a sequence consisting essentially of  $R^3 R^4 R^5 R^6 R^7 R^8$  in general formula I.
- 19. A method according to Claim 18, wherein the peptide has a sequence consisting essentially of Val Tyr Ile His Pro Phe (SEQ ID NO:3).
- 20. A method according to Claim 1, wherein the peptide has a sequence consisting essentially of  $R^5 R^6 R^7 R^8$  in general formula I.
- 21. A method according to Claim 20, wherein the peptide has a sequence consisting essentially of Ile His Pro Phe (SEQ ID NO:7).

- 22. A method according to Claim 1, wherein the peptide comprises poly-Gly.
- 23. A method according to Claim 22, wherein the peptide has a sequence selected from Asp-Arg-Val-Gly-Gly-Gly-Gly (SEQ ID NO:16) and Gly-Gly-Gly-Asp-Arg-Val (SEQ ID NO:17).

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- 24. A method according to Claim 1, wherein the peptide comprises poly-Lys.
- 25. A method according to Claim 24, wherein the peptide has a sequence selected from Arg-Val-Tyr-Ile-His-Pro-Lys-Lys-Lys (SEQ ID NO:18) and Lys-Lys-Lys-Lys-Lys-Lys-Arg-Val-Tyr-Ile-His-Pro (SEQ ID NO:19)
- 26. A method according to Claim 1, wherein the peptide comprises a D-amino acid.
- 27. A method according to Claim 26, wherein the peptide has a sequence selected from D-Ala-Ile-His-Pro-Phe (SEQ ID NO:20); and Ile-His-Pro-Phe-D-Ala (SEQ ID NO:21)
  - 28. A method according to Claim 1, wherein the peptide is PEGylated.
- 29. A method according to Claim 28, wherein the PEGylated peptide is selected from the peptides of SEQ ID NOS. 3, 5, 6, 7, 8, 9, 10 and 11.